WEST Search History

DATE: Monday, November 25, 2002

| Set Name side by side | | Hit Count | Set Name result set |
|--------------------------|---|-----------|------------------------|
| DB=US OP=ADJ | SPT,PGPB,JPAB,EPAB,DWPI; THES=ASSIGNEE; PLUR=YES; | | |
| L2 | dual specificity phosphatase same humen and (nucleic acid or gene or dna or cdna) | 40 | L2 |
| L1 | duel specificity phosphatase same humen and (nucleic acid or gene or dna or cdna) | 0 | L1 |

END OF SEARCH HISTORY

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Search Results - Record(s) 1 through 20 of 40 returned.

1. Document ID: US 20020155505 A1

L2: Entry 1 of 40

File: PGPB

Oct 24, 2002

PGPUB-DOCUMENT-NUMBER: 20020155505

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020155505 A1

TITLE: Methods for ligand discovery

PUBLICATION-DATE: October 24, 2002

INVENTOR - INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Wells, Jim Burlingame CA US
Erlanson, Dan San Francisco CA US
Braisted, Andrew C. San Francisco CA US

US-CL-CURRENT: 435/7.1; 530/324, 564/161, 564/192, 564/30, 564/84

| Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KWIC |
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2. Document ID: US 20020151007 A1

L2: Entry 2 of 40

File: PGPB

Oct 17, 2002

PGPUB-DOCUMENT-NUMBER: 20020151007

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020151007 A1

TITLE: Methods of use of a novel lysyl oxidase-related protein

PUBLICATION-DATE: October 17, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Khodadoust, Mehran M. Brookline MA US MacBeth, Kyle J. Boston MA US

US-CL-CURRENT: $\underline{435}/\underline{183}$; $\underline{435}/\underline{320.1}$, $\underline{435}/\underline{325}$, $\underline{435}/\underline{69.1}$, $\underline{536}/\underline{23.2}$

| Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KWIC |
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3. Document ID: US 20020150954 A1

Record List Display

L2: Entry 3 of 40

File: PGPB

Oct 17, 2002

PGPUB-DOCUMENT-NUMBER: 20020150954

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020150954 A1

7

TITLE: Compositions and methods for identifying agents which modulate PTEN function and

PI-3 kinase pathways

PUBLICATION-DATE: October 17, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Durden, Donald L.

Indianapolis

IN

US

US-CL-CURRENT: 435/7.23; 514/12, 514/152, 514/27, 514/283, 514/449

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KWC |
Draw, Desc | Image |

4. Document ID: US 20020137170 A1

L2: Entry 4 of 40

File: PGPB

Sep 26, 2002

PGPUB-DOCUMENT-NUMBER: 20020137170

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020137170 A1

TITLE: DSP-16 dual-specificity phosphatase

PUBLICATION-DATE: September 26, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Luche, Ralf M.

Seattle

WA

US

Wei, Bo

Kirkland

WA

US

US-CL-CURRENT: 435/196; 435/320.1, 435/325, 435/69.1, 536/23.2

Full Title Citation Front Review Classification Date Reference Sequences Attachments

Draw, Desc Image

KMC

5. Document ID: US 20020123464 A1

L2: Entry 5 of 40

File: PGPB

Sep 5, 2002

PGPUB-DOCUMENT-NUMBER: 20020123464

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020123464 A1

TITLE: 69087, 15821, and 15418, methods and compositions of human proteins and uses

thereof

PUBLICATION-DATE: September 5, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Kapeller-Libermann, Rosana Chestnut Hill MA US Bandaru, Rajasekhar Watertown MA US

US-CL-CURRENT: 514/12; 435/320.1, 435/325, 435/69.1, 530/350, 536/23.5

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | KWC |
Draw, Desc | Image |

6. Document ID: US 20020102693 A1

L2: Entry 6 of 40 File: PGPB Aug 1, 2002

PGPUB-DOCUMENT-NUMBER: 20020102693

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020102693 A1

TITLE: DSP-14 dual-specificity phosphatase

PUBLICATION-DATE: August 1, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Luche, Ralf M. Seattle WA US

US-CL-CURRENT: 435/196; 435/320.1, 435/325, 435/69.1, 536/23.2

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | KMC | Draw, Desc | Image |

7. Document ID: US 20020102691 A1

L2: Entry 7 of 40 File: PGPB Aug 1, 2002

PGPUB-DOCUMENT-NUMBER: 20020102691

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020102691 A1

TITLE: Cytokine-, stress-, and oncoprotein-activated human protein kinase kinases

PUBLICATION-DATE: August 1, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Davis, Roger J. Princeton MA US
Raingeaud, Joel Palaiseau FR
Derijard, Benoit Nice FR

US-CL-CURRENT: 435/194; 435/320.1, 435/325, 435/6, 435/69.1, 536/23.2

Full Titte Citation Front Review Classification Date Reference Sequences Attachments | KWIC | Draws Description | Image |

8. Document ID: US 20020094561 A1

L2: Entry 8 of 40

File: PGPB

Jul 18, 2002

PGPUB-DOCUMENT-NUMBER: 20020094561

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020094561 A1

TITLE: Isolated human phosphatase proteins, nucleic acid molecules encoding human

phosphatase proteins, and uses thereof

PUBLICATION-DATE: July 18, 2002

INVENTOR-INFORMATION:

| NAME | CITY | STATE | COUNTRY | RULE-47 |
|-------------------------|------------|-------|---------|---------|
| Ye, Jane | Boyds | MD | US | |
| Yan, Chunhua | Boyds | MD | US | |
| Di Francesco, Valentina | Rockville | MD | US | |
| Beasley, Ellen M. | Darnestown | MD | US | |

US-CL-CURRENT: 435/196; 435/325, 435/6, 435/69.1, 435/7.1, 536/23.2, 800/8

| Full Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | KWIC |
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| Draw, Desc III | mage | | | | | | | | |
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9. Document ID: US 20020090703 A1

L2: Entry 9 of 40

File: PGPB

Jul 11, 2002

PGPUB-DOCUMENT-NUMBER: 20020090703

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020090703 A1

TITLE: Mammalian protein phosphatases

PUBLICATION-DATE: July 11, 2002

INVENTOR - INFORMATION:

| NAME | CITY | STATE | COUNTRY | RULE-47 |
|---------------------|---------------|-------|---------|---------|
| Plowman, Gregory D. | San Carlos | CA | US | |
| Martinez, Ricardo | Foster City | CA | US | |
| Whyte, David | Belmont | CA | US | |
| Manning, Gerard | Menlo Park | CA | US | |
| Sudarsanam, Sucha | Greenbrae | CA | US | |
| Caenepeel, Sean | Oakland | CA | US | |
| Hill, Ron | Burlingame | CA | US | |
| Flanagan, Peter | San Francisco | CA | US | |

US-CL-CURRENT: 435/196; 435/320.1, 435/325, 435/69.1, 536/23.2

| Full Title Citation Front Review Classification Date Reference Sequences Attachments | KWIC |
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10. Document ID: US 20020090624 A1

Record List Display

L2: Entry 10 of 40

File: PGPB

Jul 11, 2002

PGPUB-DOCUMENT-NUMBER: 20020090624

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020090624 A1

TITLE: Gene markers useful for detecting skin damage in response to ultraviolet

radiation

PUBLICATION-DATE: July 11, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Blumenberg, Miroslav

New York

NY

US

US-CL-CURRENT: 435/6

Title Citation Front Review Classification Date Reference Sequences Attachments Draw, Desc | Image

KWIC

11. Document ID: US 20020081612 A1

L2: Entry 11 of 40

File: PGPB

Jun 27, 2002

PGPUB-DOCUMENT-NUMBER: 20020081612

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020081612 A1

TITLE: Detection and diagnosis of smoking related cancers

PUBLICATION-DATE: June 27, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Katz, Ruth

Houston

TX

US

Jiang, Feng

Houston

TX

US

US-CL-CURRENT: 435/6; 536/23.1

Title Citation Front Review Classification Date Reference Sequences Draw. Desc | Image

KWIC

12. Document ID: US 20020068287 A1

L2: Entry 12 of 40

File: PGPB

Jun 6, 2002

PGPUB-DOCUMENT-NUMBER: 20020068287

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020068287 A1

TITLE: Methods of identifying integrin ligands using differential gene expression

PUBLICATION-DATE: June 6, 2002

INVENTOR-INFORMATION:

CITY STATE COUNTRY RULE-47 NAME Brookline US Fougerolles, Antonin de MA Carulli, John Southborough MA US Kotelianski, Victor Boston MA US CTUS Green, Cynthia D. Madison Hsu, Andro Berkley CA US

US-CL-CURRENT: 435/6; 435/91.2, 536/23.1

Full Title Citation Front Review Classification Date Reference Sequences Attachments KMC Praw Desc Image

13. Document ID: US 20020065406 A1

L2: Entry 13 of 40

File: PGPB

May 30, 2002

PGPUB-DOCUMENT-NUMBER: 20020065406

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020065406 A1

TITLE: 18221, a novel dual specificity phosphatase and uses thereof

PUBLICATION-DATE: May 30, 2002

INVENTOR - INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Meyers, Rachel A.

Newton

MA

US

US-CL-CURRENT: 536/23.1; 435/196, 435/6

Full Title Citation Front Review Classification Date Reference Sequences Attachments

Draw Desc Image

KOMC

14. Document ID: US 20020034807 A1

L2: Entry 14 of 40

File: PGPB

Mar 21, 2002

PGPUB-DOCUMENT-NUMBER: 20020034807

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020034807 A1

TITLE: 38692 and 21117, novel dual specificity phosphatase molecules and uses therefor

PUBLICATION-DATE: March 21, 2002

INVENTOR - INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Meyers, Rachel A.

Newton

MA

US

US-CL-CURRENT: 435/196; 435/325, 435/6, 435/69.1, 435/7.1, 514/44, 530/388.1, 536/23.2

Full Title Citation Front Review Classification Date Reference Sequences Attachments
Draw Desc Image

KMMC

15. Document ID: US 20020009797 A1

L2: Entry 15 of 40

File: PGPB

Jan 24, 2002

PGPUB-DOCUMENT-NUMBER: 20020009797

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020009797 A1

TITLE: Growth stimulation of biological cells and tissue by electromagnetic fields and

uses thereof

PUBLICATION-DATE: January 24, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE COUNTRY

RULE-47

Wolf, David A.

Houston

ΤX

US

Goodwin, Thomas J.

Friendswood

TX US

US-CL-CURRENT: 435/289.1; 435/173.8, 435/298.2



KWIC

16. Document ID: US 20020009730 A1

L2: Entry 16 of 40

File: PGPB

Jan 24, 2002

PGPUB-DOCUMENT-NUMBER: 20020009730

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020009730 A1

TITLE: Human stress array

PUBLICATION-DATE: January 24, 2002

INVENTOR-INFORMATION:

Lukashev, Matvey E.

Draw, Desc Image

NAME

CITY

Newton

STATE

COUNTRY

RULE-47

Chenchik, Alex

Palo Alto

CA MA US US

US-CL-CURRENT: 435/6; 536/24.3

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments

MANAGE

17. Document ID: US 20010049358 A1

L2: Entry 17 of 40

File: PGPB

Dec 6, 2001

PGPUB-DOCUMENT-NUMBER: 20010049358

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20010049358 A1

TITLE: DSP-12 and DSP-13 dual-specificity phosphatases

PUBLICATION-DATE: December 6, 2001

Record List Display

INVENTOR - INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Luche, Ralf M. Wei, Bo

Seattle Kirkland WA WA US US

US-CL-CURRENT: 514/12; 435/196, 435/325, 435/6, 435/69.1, 435/7.1

Full Title Citation Front Review Classification Date Reference Sequences Attachments

Draw Desc Image

KWC

18. Document ID: US 6436685 B1

L2: Entry 18 of 40

File: USPT

Aug 20, 2002

US-PAT-NO: 6436685

DOCUMENT-IDENTIFIER: US 6436685 B1

TITLE: CSAPTP protein molecules and uses therefor

DATE-ISSUED: August 20, 2002

INVENTOR - INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Acton; Susan L.

Lexington

MA

US-CL-CURRENT: 435/196; 435/252.3, 435/254.11, 435/320.1, 435/6, 536/23.2

ABSTRACT:

The invention provides isolated <u>nucleic acid</u> molecules, designated CSAPTP <u>nucleic acid</u> molecules, which encode novel protein tyrosine phosphatases. The invention also provides antisense <u>nucleic acid</u> molecules, recombinant expression vectors containing CSAPTP <u>nucleic acid</u> molecules, host cells into which the expression vectors have been introduced, and methods for producing CSAPTP polypeptides.

15 Claims, 10 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 10

Full Title Citation Front Review Classification Date Reference Sequences Attachments

Draw, Desc Image

KWAC

19. Document ID: US 6420153 B1

L2: Entry 19 of 40

File: USPT

Jul 16, 2002

US-PAT-NO: 6420153

DOCUMENT-IDENTIFIER: US 6420153 B1

TITLE: 18232, a novel dual specificity phosphatase and uses therefor

DATE-ISSUED: July 16, 2002

INVENTOR - INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Meyers; Rachel A.

Newton

MA

Weich; Nadine

Brookline

MA

 $\text{US-CL-CURRENT: } \underline{435/196}; \ \underline{435/252.3}, \ \underline{435/320.1}, \ \underline{435/325}, \ \underline{536/23.1}, \ \underline{536/23.2}, \ \underline{536/24.1}$

ABSTRACT:

The invention provides isolated <u>nucleic acids</u> molecules, designated 18232 <u>nucleic acid</u> molecules, which encode novel dual specificity phosphatase family members. The invention also provides antisense <u>nucleic acid</u> molecules, recombinant expression vectors containing 18232 <u>nucleic acid</u> molecules, host cells into which the expression vectors have been introduced, and nonhuman transgenic animals in which a 18232 <u>gene</u> has been introduced or disrupted. The invention still further provides isolated 18232 proteins, fusion proteins, antigenic peptides and anti-18232 antibodies. Diagnostic methods utilizing compositions of the invention are also provided. The invention also provides methods of modulating the differentiation and proliferation of hematopoietic cells (e.g., erythroid cells) utilizing the compositions of the invention. Accordingly, methods of treating, preventing and/or diagnosing erythroid-associated disorders such as anemias, leukemias, and erythrocytosis are disclosed.

15 Claims, 9 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 8

Full Title Citation Front Review Classification Date Reference Sequences Attachments

Draw, Descriptings

KWC

20. Document ID: US 6335170 B1

L2: Entry 20 of 40

File: USPT

Jan 1, 2002

US-PAT-NO: 6335170

DOCUMENT-IDENTIFIER: US 6335170 B1

TITLE: Gene expression in bladder tumors

DATE-ISSUED: January 1, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Orntoft; Torben F.

DK 8230 Aabyhoj

DK

US-CL-CURRENT: 435/6; 435/91.1, 435/91.2, 536/23.1, 536/24.3, 536/24.31, 536/24.33

ABSTRACT:

Methods for analyzing tumor cells, particularly bladder tumor cells employ gene expression analysis of samples. Gene expression patterns are formed and compared to reference patterns. Alternatively gene expression patterns are manipulated to exclude genes which are expressed in contaminating cell populations. Another alternative employs subtraction of the expression of genes which are expressed in contaminating cell types. These methods provide improved accuracy as well as alternative basis for analysis from diagnostic and prognostic tools currently available.

21 Claims, 24 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 15

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Search Results - Record(s) 21 through 40 of 40 returned.

21. Document ID: US 6331614 B1

L2: Entry 21 of 40

File: USPT

Dec 18, 2001

US-PAT-NO: 6331614

DOCUMENT-IDENTIFIER: US 6331614 B1

TITLE: Human CDC14A gene

DATE-ISSUED: December 18, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Wong; Alexander K. C. La Jolla CA
Teng; David H. -F. Salt Lake City UT
Tavtigian; Sean V. Salt Lake City UT

US-CL-CURRENT: <u>536/23.5</u>; <u>435/320.1</u>, <u>435/325</u>, <u>536/23.1</u>

ABSTRACT:

The present invention relates generally to the field of human genetics. Specifically, the present invention relates to human CDC14A gene which has been found to be mutated in certain tumor cell lines. More specifically, the invention relates to a novel sequence for the human CDC14A gene. The present invention further relates to somatic mutations in the CDC14A gene in human cancer and their use in the diagnosis and prognosis of human cancer. The invention also relates to the therapy of human cancers which have a mutation in the CDC14A gene, including gene therapy, protein replacement therapy and protein mimetics. The invention further relates to the screening of drugs for cancer therapy. Finally, the invention relates to the screening of the CDC14A gene for mutations, which are useful for diagnosing the predisposition to cancer.

14 Claims, 1 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 1

Full Title Citation Front Review Classification Date Reference Sequences Attachments

Draw Desc Image

KWC

22. Document ID: US 6331396 B1

L2: Entry 22 of 40

File: USPT

Dec 18, 2001

US-PAT-NO: 6331396

DOCUMENT-IDENTIFIER: US 6331396 B1

TITLE: Arrays for identifying agents which mimic or inhibit the activity of interferons

DATE-ISSUED: December 18, 2001

INVENTOR - INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Silverman; Robert H. Beachwood OH
Williams; Bryan R. G. Cleveland OH
Der; Sandy Cleveland OH

US-CL-CURRENT: 435/6; 435/287.2, 536/23.1, 536/23.52, 536/24.3, 536/24.31

ABSTRACT:

Methods and model systems for identifying and characterizing new therapeutic agents, particularly proteins, which mimic or inhibit the activity of all interferons, Type I interferons, IFN-.alpha., IFN-.beta., or IFN-.gamma. The method comprises administering an interferon selected from the group consisting of IFN-.alpha., IFN .beta., IFN-.tau., IFN-.omega., IFN-.gamma., and combinations thereof to cultured cells, administering the candidate agent to a duplicate culture of cells; and measuring the effect of the candidate agent and the interferon on the transcription or translation of one or, preferably, a plurality of the interferon stimulated genes or the interferon repressed genes (hereinafter referred to as "ISG's" and "IRGs", respectively). The model system is an array with gene probes that hybridize with from about 100 to about 5000 ISG and IRG transcripts.

8 Claims, 0 Drawing figures Exemplary Claim Number: 1

Full Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Draw, Desc | Image |

KWIC

23. Document ID: US 6300092 B1

L2: Entry 23 of 40

File: USPT

Oct 9, 2001

US-PAT-NO: 6300092

DOCUMENT-IDENTIFIER: US 6300092 B1

TITLE: Methods of use of a novel lysyl oxidase-related protein

DATE-ISSUED: October 9, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Khodadoust; Mehran M. Brookline MA MacBeth; Kyle J. Boston MA

US-CL-CURRENT: 435/16; 435/189, 435/193, 435/25, 435/7.1

ABSTRACT:

Novel Lor-2 polypeptides, proteins, and <u>nucleic acid</u> molecules are disclosed. In addition to isolated, full-length Lor-2 proteins, the invention further provides isolated Lor-2 fusion proteins, antigenic peptides and anti-Lor-2 antibodies. The invention also provides Lor-2 <u>nucleic acid</u> molecules, recombinant expression vectors containing a <u>nucleic acid</u> molecule of the invention, host cells into which the expression vectors have been introduced and non-human transgenic animals in which a Lor-2 <u>gene</u> has been introduced or disrupted. Diagnostic, screening and therapeutic methods utilizing compositions of the invention are also provided.

17 Claims, 14 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 14 Full Title Citation Front Review Classification Date Reference Sequences Attachments KMC
Draw Desc Image

24. Document ID: US 6268135 B1

L2: Entry 24 of 40

File: USPT

Jul 31, 2001

US-PAT-NO: 6268135

DOCUMENT-IDENTIFIER: US 6268135 B1

TITLE: Phospholipase molecule and uses therefor

DATE-ISSUED: July 31, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Acton; Susan Lexington MA

US-CL-CURRENT: $\underline{435}/\underline{6}$; $\underline{435}/\underline{198}$, $\underline{435}/\underline{21}$, $\underline{435}/\underline{252.3}$, $\underline{435}/\underline{320.1}$, $\underline{530}/\underline{350}$, $\underline{536}/\underline{23.2}$,

<u>536/23.5</u>

ABSTRACT:

Novel CSAPL polypeptides, proteins, and <u>nucleic acid</u> molecules are disclosed. In addition to isolated, full-length CSAPL proteins, the invention further provides isolated CSAPL fusion proteins, antigenic peptides and anti-CSAPL antibodies. The invention also provides CSAPL <u>nucleic acid</u> molecules, recombinant expression vectors containing a <u>nucleic acid</u> molecule of the invention, host cells into which the expression vectors have been introduced and non-human transgenic animals in which a CSAPL <u>gene</u> has been introduced or disrupted. Diagnostic, screening and therapeutic methods utilizing compositions of the invention are also provided.

14 Claims, 3 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 3

| Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments |
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KWIC

25. Document ID: US 6258582 B1

L2: Entry 25 of 40

File: USPT

Jul 10, 2001

US-PAT-NO: 6258582

DOCUMENT-IDENTIFIER: US 6258582 B1

TITLE: CSAPTP nucleic acid molecules and uses therefor

DATE-ISSUED: July 10, 2001

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Acton; Susan Jamaica Plain MA

US-CL-CURRENT: 435/196; 435/252.3, 435/320.1, 435/69.1, 530/350, 536/23.2, 536/23.5

ABSTRACT:

Novel CSAPTP polypeptides, proteins, and nucleic acid molecules are disclosed. In addition to isolated, full-length CSAPTP proteins, the invention further provides isolated CSAPTP fusion proteins, antigenic peptides and anti-CSAPTP antibodies. The invention also provides CSAPTP <u>nucleic acid</u> molecules, recombinant expression vectors containing a <u>nucleic acid</u> molecule of the invention, host cells into which the expression vectors have been introduced and non-human transgenic animals in which a CSAPTP gene has been introduced or disrupted. Diagnostic, screening and therapeutic methods utilizing compositions of the invention are also provided.

23 Claims, 10 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 10

Title Citation Front Review Classification Date Reference Sequences Attachments

KWAC

26. Document ID: US 6174676 B1

L2: Entry 26 of 40

File: USPT

Jan 16, 2001

US-PAT-NO: 6174676

DOCUMENT-IDENTIFIER: US 6174676 B1

TITLE: Cytokine-stress- and oncoprotein-activated human protein kinase kinases

DATE-ISSUED: January 16, 2001

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Davis; Roger J.

Princeton

FR

Raingeaud; Joel

Palaiseau

FR

Derijard; Benoit

Nice

US-CL-CURRENT: $\underline{435/6}$; $\underline{435/194}$, $\underline{435/810}$, $\underline{435/975}$, $\underline{436/501}$, $\underline{436/94}$, $\underline{530/387.1}$, $\underline{536/23.1}$, 536/24.3, 536/24.33, 536/25.3

ABSTRACT:

Disclosed are human mitogen-activated (MAP) kinase kinase isoforms (MKKs). MKKs mediate unique signal transduction pathways that activate human-MAP kinases p38 and JNK, which result in activation of other factors, including activating transcription factor-2 (ATF2) and c-Jun. The pathways are activated by a number of factors, including cytokines and environmental stress. Methods are provided for identifying reagents that modulate MKK function or activity and for the use of such reagents in the treatment of MKK-mediated disorders.

29 Claims, 28 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 28

> Full Title Citation Front Review Classification Date Reference Sequences Attachments Draw, Desc Image

KWIC

27. Document ID: US 6162897 A

L2: Entry 27 of 40

File: USPT

Dec 19, 2000

US-PAT-NO: 6162897

DOCUMENT-IDENTIFIER: US 6162897 A

TITLE: 17q-linked breast and ovarian cancer susceptibility gene

DATE-ISSUED: December 19, 2000

INVENTOR - INFORMATION:

| NAME | CITY | STATE | ZIP CODE | COUNTRY |
|---------------------------|----------------|-------|----------|---------|
| Skolnick; Mark H. | Salt Lake City | UT | | |
| Goldgar; David E. | Salt Lake City | UT | | |
| Miki; Yoshio | Salt Lake City | UT | | |
| Swenson; Jeff | Salt Lake City | UT | | |
| Kamb; Alexander | Salt Lake City | UT | | • |
| Harshman; Keith D. | Salt Lake City | UT | | |
| Shattuck-Eidens; Donna M. | Salt Lake City | UT | | |
| Tavtigian; Sean V. | Salt Lake City | UT | | |
| Wiseman; Roger W. | Durham | NC | | |
| Futreal; P. Andrew | Durham | NC | | |

US-CL-CURRENT: <u>530/350</u>; <u>424/174.1</u>, <u>435/7.1</u>

ABSTRACT:

The present invention relates generally to the field of human genetics. Specifically, the present invention relates to methods and materials used to isolate and detect a human breast and ovarian cancer predisposing gene (BRCA1), some mutant alleles of which cause susceptibility to cancer, in particular breast and ovarian cancer. More specifically, the invention relates to germline mutations in the BRCA1 gene and their use in the diagnosis of predisposition to breast and ovarian cancer. The present invention further relates to somatic mutations in the BRCA1 gene in human breast and ovarian cancer and their use in the diagnosis and prognosis of human breast and ovarian cancer. Additionally, the invention relates to somatic mutations in the BRCA1 gene in other human cancers and their use in the diagnosis and prognosis of human cancers. The invention also relates to the therapy of human cancers which have a mutation in the BRCA1 gene, including gene therapy, protein replacement therapy and protein mimetics. The invention further relates to the screening of drugs for cancer therapy. Finally, the invention relates to the screening of the BRCA1 gene for mutations, which are useful for diagnosing the predisposition to breast and ovarian cancer.

3 Claims, 19 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 18

| Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments |
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| Drawi D | eso Ir | nage | | | | | | | |

KMC

28. Document ID: US 6136596 A

L2: Entry 28 of 40

File: USPT

Oct. 24, 2000

US-PAT-NO: 6136596

DOCUMENT-IDENTIFIER: US 6136596 A

TITLE: Cytokine-, stress-, and oncoprotein-activated human protein kinase kinases

DATE-ISSUED: October 24, 2000

INVENTOR - INFORMATION:

NAME

CITY

STATE

ZIP CODE

COUNTRY

Davis; Roger J. Whitmarsh; Alan

Princeton Shrewsbury MA MA

Tournier; Cathy

Worcester

MA

US-CL-CURRENT: <u>435/325</u>; <u>435/194</u>, <u>435/252.3</u>, <u>435/320.1</u>, <u>536/23.2</u>

ABSTRACT:

Disclosed are human mitogen-activated (MAP) kinase kinase isoforms (MKKs). MKKs mediate unique signal transduction pathways that activate human MAP kinases p38 and JNK, which result in activation of other factors, including activating transcription factor-2 (ATF2) and c-Jun. The pathways are activated by a number of factors, including cytokines and environmental stress. Methods are provided for identifying reagents that modulate MKK function or activity and for the use of such reagents in the treatment of MKK-mediated disorders.

9 Claims, 54 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 54

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments |
Draw, Desc | Image |

KWIC

29. Document ID: US 6074851 A

L2: Entry 29 of 40

File: USPT

Jun 13, 2000

US-PAT-NO: 6074851

DOCUMENT-IDENTIFIER: US 6074851 A

TITLE: Catalytic macro molecules having cdc25B like activity

DATE-ISSUED: June 13, 2000

INVENTOR-INFORMATION:

NAME

CITY

STATE

ZIP CODE COUNTRY

Deibel, Jr.; Martin R.

Kalamazoo

MI .

Yem; Anthony W. Wolfe; Cindy L.

Kalamazoo

MI

Portage

MI

US-CL-CURRENT: 435/69.7; 435/194

ABSTRACT:

This invention discloses novel forms of catalytic macro molecules that are related to cdc25B, a cell cycle specific phosphatase. These special domains of cdc25B, special fusions with GST, and unique peptides and proteins, their utility, and the method of making them are all described.

3 Claims, 6 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 9

Full Title Citation Front Review Classification Date Reference Sequences Attachments
Draw. Desc Image

KWC

30. Document ID: US 5998188 A

L2: Entry 30 of 40

File: USPT

Dec 7, 1999

US-PAT-NO: 5998188

DOCUMENT-IDENTIFIER: US 5998188 A

TITLE: Mitogen activated protein kinase phosphatase <u>cDNAS</u> and their biologically active

expression products

DATE-ISSUED: December 7, 1999

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Stork; Philip J. S.

Portland

OR OR

Misra-Press; Anita Portland

US-CL-CURRENT: 435/196

ABSTRACT:

The invention relates to a novel mitogen-activated protein kinase phosphatase, MKP-2. The invention further relates to methods and means for preparing and to <u>nucleic acids</u> encoding this protein. The MKP-2 of the present invention is useful in the control of cell growth, differentiation and apoptosis.

3 Claims, 44 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 24

| Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments |
|---------|-------|----------|-------|--------|----------------|------|-----------|-----------|-------------|
| Draw, D | esc l | mage | | | | | | | |

KWIC

31. Document ID: US 5804427 A

L2: Entry 31 of 40

File: USPT

Sep 8, 1998

US-PAT-NO: 5804427

DOCUMENT-IDENTIFIER: US 5804427 A

TITLE: Cytokine-, stress-, and oncoprotein-activated human protein kinase kinases

DATE-ISSUED: September 8, 1998

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Davis; Roger Princeton MA

Raingeaud; Joel Bazoges en Pareds FR
Derijard; Benoit Marseilles FR

US-CL-CURRENT: 435/194; 435/183, 530/350

ABSTRACT:

Disclosed are human mitogen-activated (MAP) kinase kinase isoforms (MKKs). MKKs mediate unique signal transduction pathways that activate human MAP kinases p38 and JNK, which result in activation of other factors, including activating transcription factor-2 (ATF2) and c-Jun. The pathways are activated by a number of factors, including

cytokines and environmental stress. Methods are provided for identifying reagents that modulate MKK function or activity and for the use of such reagents in the treatment of MKK-mediated disorders.

4 Claims, 3 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 3

Full Title Citation Front Review Classification Date Reference Sequences Attachments

Draw Desc Image

32. Document ID: US 5753441 A

L2: Entry 32 of 40

File: USPT

May 19, 1998

US-PAT-NO: 5753441

DOCUMENT-IDENTIFIER: US 5753441 A

TITLE: 170-linked breast and ovarian cancer susceptibility gene

DATE-ISSUED: May 19, 1998

INVENTOR - INFORMATION:

| NAME | CITY | STATE | ZIP CODE | COUNTRY |
|---------------------------|----------------|-------|----------|---------|
| Skolnick; Mark H. | Salt Lake City | UT | | |
| Goldgar; David E. | Salt Lake City | UT | | |
| Miki; Yoshio | Salt Lake City | UT | | |
| Swenson; Jeff | Salt Lake City | UT | | |
| Kamb; Alexander | Salt Lake City | UT | | |
| Harshman; Keith D. | Salt Lake City | UT | | |
| Shattuck-Eidens; Donna M. | Salt Lake City | UT | | |
| Tavtigian; Sean V. | Salt Lake City | UT | | |
| Wiseman; Roger W. | Durham | NC | | |
| Futreal; P. Andrew | Durham | NC | | |

 $\begin{array}{l} \text{US-CL-CURRENT: } \underline{435/6}; \ \underline{424/1.11}, \ \underline{435/4}, \ \underline{435/7.1}, \ \underline{435/7.2}, \ \underline{435/7.9}, \ \underline{435/91.1}, \ \underline{435/91.2}, \\ \underline{436/500}, \ \underline{436/548}, \ \underline{530/387.2}, \ \underline{530/388.1}, \ \underline{536/23.1}, \ \underline{536/24.3}, \ \underline{536/24.33} \end{array}$

ABSTRACT:

The present invention relates generally to the field of human genetics. Specifically, the present invention relates to methods and materials used to isolate and detect a human breast and ovarian cancer predisposing gene (BRCA1), some mutant alleles of which cause susceptibility to cancer, in particular breast and ovarian cancer. More specifically, the invention relates to germline mutations in the BRCA1 gene and their use in the diagnosis of predisposition to breast and ovarian cancer. The present invention further relates to somatic mutations in the BRCA1 gene in human breast and ovarian cancer and their use in the diagnosis and prognosis of human breast and ovarian cancer. Additionally, the invention relates to somatic mutations in the BRCA1 gene in other human cancers and their use in the diagnosis and prognosis of human cancers. The invention also relates to the therapy of human cancers which have a mutation in the BRCA1 gene, including gene therapy, protein replacement therapy and protein mimetics. The invention further relates to the screening of drugs for cancer therapy. Finally, the invention relates to the screening of the BRCA1 gene for mutations, which are useful for diagnosing the predisposition to breast and ovarian cancer.

37 Claims, 19 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 18 Full Title Citation Front Review Classification Date Reference Sequences Attachments KWC Draw, Description

33. Document ID: US 5736381 A

L2: Entry 33 of 40

File: USPT

Apr 7, 1998

US-PAT-NO: 5736381

DOCUMENT-IDENTIFIER: US 5736381 A

TITLE: Cytokine-, stress-, and oncoprotein-activated human protein kinase kinases

DATE-ISSUED: April 7, 1998

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Davis; Roger J. Princeton MA 01541

Gupta; Shashi Worcester MA 01604

Raingeaud; Joel 85390 Bazoges en Pareds FR
Derijard; Benoit 13012 Marseille FR

US-CL-CURRENT: $\frac{435}{252.3}$; $\frac{435}{320.1}$, $\frac{435}{325}$, $\frac{435}{6}$, $\frac{435}{69.1}$, $\frac{435}{91.1}$, $\frac{536}{23.1}$, $\frac{536}{23.5}$, $\frac{536}{24.31}$, $\frac{536}{24.33}$

ABSTRACT:

Disclosed are human mitogen-activated (MAP) kinase kinase isoforms (MKKs). MKKs mediate unique signal transduction pathways that activate human MAP kinases p38 and JNK, which result in activation of other factors, including activating transcription factor-2 (ATF2) and c-Jun. The pathways are activated by a number of factors, including cytokines and environmental stress. Methods are provided for identifying reagents that modulate MKK function or activity and for the use of such reagents in the treatment of MKK-mediated disorders.

20 Claims, 8 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 28

| Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments |
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| Draw, D | eso Ir | mage | | | | -1 | | | |

KWIC

34. Document ID: US 5710001 A

L2: Entry 34 of 40

File: USPT

Jan 20, 1998

US-PAT-NO: 5710001

DOCUMENT-IDENTIFIER: US 5710001 A

TITLE: 17q-linked breast and ovarian cancer susceptibility gene

DATE-ISSUED: January 20, 1998

INVENTOR - INFORMATION:

| NAME | CITY | STATE | ZIP CODE | COUNTRY |
|---------------------------|----------------|-------|----------|---------|
| Skolnick; Mark H. | Salt Lake City | UT | | |
| Goldgar; David E. | Salt Lake City | UT | | |
| Miki; Yoshio | Salt Lake City | UT | | |
| Swenson; Jeff | Salt Lake City | UT | | |
| Kamb; Alexander | Salt Lake City | UT | | |
| Harshman; Keith D. | Salt Lake City | UT | | |
| Shattuck-Eidens; Donna M. | Salt Lake City | UT | | |
| Tavtigian; Sean V. | Salt Lake City | UT | | |
| Wiseman; Roger W. | Durham | °NC | | |
| Futreal; P. Andrew | Durham | NC | | |

US-CL-CURRENT: $\underline{435}/\underline{6}$; $\underline{435}/\underline{7.1}$, $\underline{435}/\underline{7.9}$, $\underline{435}/\underline{91.2}$, $\underline{530}/\underline{300}$, $\underline{530}/\underline{350}$, $\underline{530}/\underline{388.1}$, $\underline{536}/\underline{24.3}$, $\underline{536}/\underline{24.33}$

ABSTRACT:

The present invention relates generally to the field of human genetics. Specifically, the present invention relates to methods and materials used to isolate and detect a human breast and ovarian cancer predisposing gene (BRCA1), some mutant alleles of which cause susceptibility to cancer, in particular breast and ovarian cancer. More specifically, the invention relates to germline mutations in the BRCA1 gene and their use in the diagnosis of predisposition to breast and ovarian cancer. The present invention further relates to somatic mutations in the BRCA1 gene in human breast and ovarian cancer and their use in the diagnosis and prognosis of human breast and ovarian cancer. Additionally, the invention relates to somatic mutations in the BRCA1 gene in other human cancers and their use in the diagnosis and prognosis of human cancers. The invention also relates to the therapy of human cancers which have a mutation in the BRCA1 gene, including gene therapy, protein replacement therapy and protein mimetics. The invention further relates to the screening of drugs for cancer therapy. Finally, the invention relates to the screening of the BRCA1 gene for mutations, which are useful for diagnosing the predisposition to breast and ovarian cancer.

35 Claims, 19 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 18

35. Document ID: US 5709999 A

L2: Entry 35 of 40

File: USPT

Jan 20, 1998

US-PAT-NO: 5709999

DOCUMENT-IDENTIFIER: US 5709999 A

TITLE: Linked breast and ovarian cancer susceptibility gene

DATE-ISSUED: January 20, 1998

INVENTOR-INFORMATION:

| NAME | CITY | STATE | ZIP | CODE | COUNTRY |
|---------------------------|---------------------------|-------|-----|------|---------|
| Shattuck-Eidens; Donna M. | Salt Lake City | UT | | | |
| Simard; Jacques | St. Augustin de Desmaures | | | | CA |
| Durocher; Francine | Ste-Foy | | | | CA |
| Emi; Mitsuuru | Tokyo | | | | JP |
| Nakamura; Yusuke | Yokohama | | | | JP |

US-CL-CURRENT: 435/6; 435/91.2, 536/23.1, 536/24.3, 536/24.33

ABSTRACT:

The present invention relates generally to the field of human genetics. Specifically, the present invention relates to methods and materials used to isolate and detect a human breast and ovarian cancer predisposing gene (BRCA1), some mutant alleles of which cause susceptibility to cancer, in particular breast and ovarian cancer. More specifically, the invention relates to germline mutations in the BRCA1 gene and their use in the diagnosis of predisposition to breast and ovarian cancer. The present invention further relates to somatic mutations in the BRCA1 gene in human breast and ovarian cancer and their use in the diagnosis and prognosis of human breast and ovarian cancer. Additionally, the invention relates to somatic mutations in the BRCA1 gene in other human cancers and their use in the diagnosis and prognosis of human cancers. The invention also relates to the therapy of human cancers which have a mutation in the BRCA1 gene, including gene therapy, protein replacement therapy and protein mimetics. The invention further relates to the screening of drugs for cancer therapy. Finally, the invention relates to the screening of the BRCA1 gene for mutations, which are useful for diagnosing the predisposition to breast and ovarian cancer.

35 Claims, 19 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 18

| Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments |
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KWIC

36. Document ID: US 5573935 A

L2: Entry 36 of 40

File: USPT

STATE

ZIP CODE

Nov 12, 1996

COUNTRY

US-PAT-NO: 5573935

DOCUMENT-IDENTIFIER: US 5573935 A

TITLE: Protein tyrosine kinase A6

DATE-ISSUED: November 12, 1996

INVENTOR-INFORMATION:

NAME CITY

Beeler; John F. Bethesda MD Larochelle; William Gaithersburg MD

Aaronson; Stuart A. Great Falls VA

US-CL-CURRENT: 435/194; 435/252.3, 435/252.33, 435/320.1, 435/69.8, 536/23.2, 536/23.5,

930/240

ABSTRACT:

A novel protein tyrosine kinase (A6) exhibiting no significant similarity to any known kinase. This protein in widely expressed throughout the body and is present in a variety of vertebrates. The $\underline{\text{cDNA}}$ was expressed in bacteria as a fusion protein which was both autophosphorylated $\underline{\text{and}}$ exhibited kinase activity toward exogenous substrates. Potential uses of this invention include immunodiagnostics and antiproliferative therapeutics.

10 Claims, 1 Drawing figures Exemplary Claim Number: 1,9 Number of Drawing Sheets: 1

| Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | KWIC |
|---------|--------|----------|-------|--------|----------------|------|-----------|-----------|-------------|------|
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37. Document ID: WO 200231132 A2 AU 200211597 A

L2: Entry 37 of 40

File: DWPI

Apr 18, 2002

DERWENT-ACC-NO: 2002-416863

DERWENT-WEEK: 200254

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TITLE: Novel <u>human dual specificity phosphatase</u>, 8843 and polynucleotides for identifying modulators for use in treating or preventing an erythroid-associated disorder e.g. anemia or leukemia in a subject

INVENTOR: WEICH, N

PRIORITY-DATA: 2000US-0686673 (October 11, 2000)

PATENT-FAMILY:

 PUB-NO
 PUB-DATE
 LANGUAGE
 PAGES
 MAIN-IPC

 WO 200231132 A2
 April 18, 2002
 E
 125
 C12N009/16

 AU 200211597 A
 April 22, 2002
 000
 C12N009/16

INT-CL (IPC): C12 N 9/16; C12 N 15/12

ABSTRACTED-PUB-NO: WO 200231132A

BASIC-ABSTRACT:

NOVELTY - An isolated <u>human dual specificity phosphatase</u> polypeptide (I), 8843, having a sequence (S1) of 201 amino acids (aa) as given in specification, or a sequence encoded by polynucleotide having a sequence of 839 or 606 bp as given in the specification or its complement, is new.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for the following:

- (1) an isolated 8843 <u>nucleic acid</u> (NA) molecule (II) encoding (I);
- (2) modulating (M1) erythropoiesis or proliferation, differentiation or survival of CD34 positive cell, by contacting an positive cell with an agent that modulates the activity or expression of (I)/(II), thereby modulates proliferation, differentiation, or survival of the cell; and
- (3) treating or preventing (M2) an erythroid-associated disorder in a subject by administering an agent that modulates activity or expression of (I).

ACTIVITY - Antianemia; Cytostatic; Nephrotrophic; Antiarthritic; Antirheumatic; Anti-human immunodeficiency virus (HIV); Dermatological. No supporting data is given.

MECHANISM OF ACTION - Gene therapy; Modulator of (I) or (II).

USE - (I) is useful for identifying a compound modulates the activity of (I). M1 is useful for modulating erythropoiesis in a subject preferably human having an erythroid associated disorder. (I) and (II) are useful for evaluating the efficacy of a treatment of an erythroid associated disorder which is an anemia or leukemia, in a subject; and for diagnosing the disorder by evaluating and comparing expression or activity of (I)/(II). M1 is useful for modulating erythropoiesis or proliferation, differentiation or survival of CD34 positive cell; and M2 is useful for treating or preventing an erythroid-associated disorder in a subject (claimed). The modulator thus identified is also useful for treating chronic renal failure, malignancies, adult and juvenile rheumatoid arthritis, disorders of hemoglobin synthesis, prematurity and zidovudine treatment of HIV infection; and disorders of liver e.g. jaundice, kidney e.g. Heymann nephritis, lung e.g. congenital anomalies and skin (dermal) e.g. seborrheic keratoses. (I) is useful for producing antibodies, in drug screening assays, in competition

binding assays to discover compounds that interact with the protein, in pharmacogenomic analysis and for monitoring therapeutic effects during clinical trials and other treatment. (IV) is useful for isolating (I), to assess abnormal tissue distribution or abnormal expression during development, to identify protein turnover, to assess normal and aberrant subcellular localization of cells in various tissues in an organism, to diagnostically monitor protein levels in tissue in pharmacogenomic analysis, for tissue typing, forensic identification, inhibiting protein function and to block ligand binding. (II) is useful as hybridization probes for $\underline{\text{cDNA}}$ and genomic $\underline{\text{DNA}}$ to isolate a full-length $\underline{\text{cDNA}}$ and genomic clones encoding (I), as primers for polymerase chain reaction (PCR) to amplify any given region of the polynucleotide, for expressing antiqenic peptides, as probes for determining the chromosomal positions of the polynucleotides, for designing ribozymes, constructing host cells, transgenic animals and for identifying a disease or disorder associated with aberrant expression or activity of (II). Fragments of (II) are also useful to synthesize antisense molecules of desired length and sequences. (II) is also useful to detect mutations in genes and gene expression products such as mRNA, as antisense constructs to control gene expression and for chromosome identification. (III) is useful for producing proteins and polypeptides, for conducting cell-based assays involving the protein or fragments and to produce non-human transgenic animals which are useful for studying the function of a receptor protein and identifying and evaluating modulators of the protein activity.

| Full | Title | Citation | Front | Classification | | Sequences | KWIC |
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38. Document ID: US 6331614 B1

L2: Entry 38 of 40

File: DWPI

Dec 18, 2001

DERWENT-ACC-NO: 2002-129551

DERWENT-WEEK: 200217

COPYRIGHT 2002 DERWENT INFORMATION LTD

TITLE: <u>Nucleic acid</u> encoding mutated form of <u>human dual-specificity phosphatase</u> CDC14A polypeptide, useful to diagnose and treat cancers

INVENTOR: TAVTIGIAN, S V; TENG, D H F; WONG, A K C

PRIORITY-DATA: 1998US-113833P (December 23, 1998), 1999US-0468872 (December 22, 1999)

PATENT-FAMILY:

 PUB-NO
 PUB-DATE
 LANGUAGE
 PAGES
 MAIN-IPC

 US 6331614 B1
 December 18, 2001
 022
 C07H021/04

INT-CL (IPC): $\underline{\text{C07}}$ $\underline{\text{H}}$ $\underline{\text{21}/\text{02}}$; $\underline{\text{C07}}$ $\underline{\text{H}}$ $\underline{\text{21}/\text{04}}$; $\underline{\text{C12}}$ $\underline{\text{N}}$ $\underline{\text{15}/\text{00}}$; $\underline{\text{C12}}$ $\underline{\text{N}}$ $\underline{\text{15}/\text{09}}$; $\underline{\text{C12}}$ $\underline{\text{N}}$ $\underline{\text{15}/\text{63}}$

ABSTRACTED-PUB-NO: US 6331614B

BASIC-ABSTRACT:

NOVELTY - An isolated $\underline{\text{nucleic acid}}$ (N1) encoding a CDC14A polypeptide (P1) with a fully defined sequence (S1) $\overline{\text{of }594\text{ amino}}$ acids as given in the specification comprising a fully defined sequence (S2) of 1785 nucleotides as given in the specification, the complement of S2 or an RNA molecule corresponding to S2, is new.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for the following:

- (1) an expression vector comprising N1 operably linked to a promoter that directs expression of the <u>nucleic acid</u>;
- (2) a vector (V1) which comprises N1;
- (3) a host cell (H1) transformed in vitro with V1; and
- (4) producing P1 comprising culturing H1 under conditions suitable for production of P1

and recovering P1.

ACTIVITY - Cytostatic.

MECHANISM OF ACTION - Gene therapy; protein replacement therapy; protein mimetics. No supporting data is given.

USE - N1 and P1 are useful to diagnose and treat human cancers which have a mutation in the CDC14A gene, by gene therapy, protein replacement therapy or protein mimetics. They can also be used to screen for drugs to treat cancer.

Full Title Citation Front Review Classification Date Reference Sequences Attachments

Draw, Desc Image

KWIC

39. Document ID: WO 200173060 A2 AU 200147782 A

L2: Entry 39 of 40

File: DWPI

Oct 4, 2001

DERWENT-ACC-NO: 2001-616517

DERWENT-WEEK: 200261

COPYRIGHT 2002 DERWENT INFORMATION LTD

TITLE: New polypeptide for modulating peptide activity and preventing hematopoietic disorders comprises the human dual specificity phosphatase 18221 protein

INVENTOR: MEYERS, R A

PRIORITY-DATA: 2000US-191858P (March 24, 2000)

PATENT-FAMILY:

 PUB-NO
 PUB-DATE
 LANGUAGE
 PAGES
 MAIN-IPC

 WO 200173060 A2
 October 4, 2001
 E
 138
 C12N015/55

 AU 200147782 A
 October 8, 2001
 000
 C12N015/55

INT-CL (IPC): A61 K 31/7088; A61 K 39/395; C07 K 16/40; C12 N 5/10; C12 N 9/16; C12 N 15/55; C12 Q 1/42; C12 Q 1/68; G01 N 33/68

ABSTRACTED-PUB-NO: WO 200173060A BASIC-ABSTRACT:

NOVELTY - An isolated $\underline{\text{nucleic acid}}$ molecule (I) encoding the $\underline{\text{human dual specificity}}$ $\underline{\text{phosphatase}}$ 18221 protein, is new.

DETAILED DESCRIPTION - An isolated <u>nucleic acid</u> (I) comprises:

- (i) a <u>nucleic acid</u> molecule comprising a nucleotide sequence at least 80% homologous to a fully defined 1292 (S1) or 654 (S2) nucleotides, or the <u>cDNA</u> insert of a plasmid (II) deposited with the American Type Culture Collection (Accession number pending);
- (ii) a <u>nucleic acid</u> molecule comprising a fragment of at least 462 nucleotides of (S1) or (S2), or a fragment of (II);
- (iii) a <u>nucleic acid</u> molecule encoding a polypeptide (III) comprising a fully defined sequence of 218 amino acids, or the amino acid sequence encoded by (II);
- (iv) a <u>nucleic acid</u> molecule encoding a fragment of (III), or a fragment of the amino acid sequence encoded by (II), where the fragment is at least 15 contiguous amino acids; and
- (v) a <u>nucleic acid</u> molecule encoding a naturally allelic variant of (III), or the amino acid encoded by (II), where the <u>nucleic acid</u> molecule hybridizes to (S1) or (S2), or a complement, under stringent conditions.

INDEPENDENT CLAIMS are also included for the following:

- (1) a host cell comprising (I);
- (2) an isolated polypeptide (III) selected from the following:
- (i) a polypeptide encoded by a <u>nucleic acid</u> molecule at least 80% homologous to (S1) or (S2), or (II);
- (ii) a naturally occurring allelic variant of (III), or the amino acid encoded by (II), where the polypeptide is encoded by (I), or its complement, under stringent conditions; and
- (iii) a fragment of at least 15 contiguous amino acids, of (III) or a polypeptide encoded by (II);
- (3) an antibody which selectively binds to (III);
- (4) producing (III), comprising culturing the host cell under conditions suitable for the expression of (I);
- (5) detecting (III) in a sample comprising contacting the sample with a compound that selectively binds to (III) and determining the binding;
- (6) a kit comprising a compound that selectively binds to (III), and instructions for use;
- (7) detecting the presence of (I) comprising contacting the sample with a <u>nucleic acid</u> probe or primer that selectively hybridizes to (I), and determining any binding;
- (8) detecting a compound that binds to (III), comprising contacting (III) or a cell expressing (III) with a test compound, and determining whether (III) binds to the test compound;
- (9) modulating (III) activity, comprising contacting (III) or a cell expressing (III) with a compound in order to modulate the activity of (III);
- (10) evaluating the efficacy of a treatment of a hematopoietic disorder in a subject, comprising:
- (i) treating a subject with a protocol under evaluation;
- (ii) assessing the expression level of a $\underline{\text{nucleic acid}}$ after the treatment, relative to the level prior to treatment; and
- (11) diagnosing or staging a hematopoietic disorder in a subject.

ACTIVITY - Cytostatic.

MECHANISM OF ACTION - Dual specificity phosphatase activity.

No biological data was provided.

USE - The polypeptides and polynucleotides are useful therapeutically to treat disorders characterized by insufficient or aberrant 18221 protein production. They can be used to evaluate the efficacy of treatment of hematopoietic disorders, especially erythroid-associated disorders (claimed), and to diagnose or stage hematopoietic disorders by detecting changes in expression relative to normal subjects. The polypeptides are useful to identify compounds binding to (claimed) and/or increasing/decreasing polypeptide activity (claimed), useful therapeutically, especially to treat or prevent hematopoietic disorders (claimed). For example, a compound modulating polypeptide activity/expression (e.g. a peptide; claimed) could be used to modulate hematopoiesis, especially in vivo in humans (claimed), by contacting a hematopoietic cell (e.g. an erythroid progenitor or differentiated cell; claimed), with the compound to alter cell proliferation, differentiation or survival (claimed). Such compounds are useful to treat conditions which involve increased hematopoietic cell activity or proliferation e.g. leukemias or decreased hematopoietic cell differentiation e.g. anemias, or to treat cancers e.g. renal carcinoma. The polypeptides can also be used to produce compounds selectively binding the polypeptide (especially antibodies), useful to detect the polypeptides (claimed) e.g. for disease

diagnosis; kits are provided (claimed). The polynucleotides can also be used to identify compounds modulating polynucleotide activity/expression (e.g. antisense molecules; claimed), useful to modulate hematopoiesis by contacting with a hematopoietic cell as above (claimed). The polynucleotides can also be used to produce nucleic acid probes/primers selectively hybridizing to the polynucleotides (e.g. nucleic acid probes hybridizing to mRNA molecules; claimed), useful to detect the polynucleotides (claimed) e.g. in disease diagnosis; kits are provided (claimed).

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments |
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L2: Entry 40 of 40

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Jul 6, 2000

DERWENT-ACC-NO: 2000-452383

DERWENT-WEEK: 200050

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TITLE: New isolated <u>nucleic acid</u> molecules encoding <u>human</u> nuclear <u>dual specificity</u> phosphatase-like protein for diagnosis of androgen independent prostate cancers

INVENTOR: RICHARDSON, J; SHYJAN, A W; VASSILIADIS, J

PRIORITY-DATA: 1998US-0223626 (December 29, 1998)

PATENT-FAMILY:

 PUB-NO
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INT-CL (IPC): <u>C12</u> <u>N</u> <u>0</u>/<u>00</u>

ABSTRACTED-PUB-NO: WO 200039277A

BASIC-ABSTRACT:

NOVELTY - Isolated <u>nucleic acid</u> molecules (I) comprise the 2860 base pair (bp) or the 2436 bp sequence provided in the specification encoding <u>human</u> nuclear <u>dual specificity</u> <u>phosphatase</u>-like protein (NDSP) or the <u>nucleic acid</u> molecule which encodes the 812 amino acid sequence provided in the specification.

DETAILED DESCRIPTION - INDEPENDENT CLAIMS are also included for the following:

- (1) a host cell containing (I);
- (2) an isolated polypeptide (II) comprising;
- (a) a fragment of the 812 amino acid sequence provided in the specification, where the fragment comprises at least 15 contiguous amino acids; or
- (b) a naturally occurring allelic variant of the 812 amino acid sequence provided in the specification, where the polypeptide is encoded by a <u>nucleic acid</u> comprising a sequence at least 55 % identical to the 2436 bp sequence provided in the specification;
- (3) an antibody which selectively binds to (II);
- (4) producing (II);
- (5) a method (M1) for detecting the presence of (II);
- (6) a kit comprising a compound which selectively binds to (II) or selectively hybridizes (I);

- (7) a method (M2) for detecting the presence of (I);
- (8) modulating the activity of (II) comprising contacting (II) or a cell expressing (II) with a compound which binds to the polypeptide;
- (9) identifying a compound which modulates the activity of (II);
- (10) a method (M3) for identifying a compound useful for treating prostate cancer comprising;
- (a) measuring the expression level of NDSP in a biological sample comprising an androgen independent prostate cancer cell; and
- (b) comparing the expression of NDSP in the presence and absence of the compound, where the compound is useful for treating prostate cancer when the expression level of NDSP in the presence of the compound is less than its expression level in the absence of the compound;
- (11) determining if a prostate cancer in a patient not undergoing androgen withdrawal therapy is androgen dependent comprising;
- (a) providing a sample of patient prostate cancer cells; and
- (b) determining if NDSP is expressed in the sample of prostate cancer cells, where the absence of NDSP expression indicates that the prostate cancer is androgen independent; and
- (12) determining the efficacy of androgen withdrawal treatment in a prostate cancer patient comprising;
- (a) providing a biological sample from the patient from a first time point and determining the expression level of NDSP;
- (b) providing a biological sample from the patient at a second time point, occurring after the patient has begun androgen withdrawal treatment and determining the expression level of NDSP; and
- (c) comparing the expression levels of NDSP in the first and second samples, where an increase in the expression level in the second sample compared with the first sample indicates that the androgen withdrawal treatment has become less effective.
- USE (I) and (II) can be used to detect androgen independent prostate cancers.

ADVANTAGE - Androgen independent prostate cancers can be distinguished from androgen dependent prostate cancers and a more suitable treatment regime can be implemented. Current treatment for prostate cancers involve androgen withdrawal, which is an ineffective treatment for androgen independent prostate cancer.

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